

PUR 4-IN-1 SKIN TINT MN4- zinc oxide lotion
PUR Minerals

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[illegible]

PUR 4-in-1 Skin Tint MN4

Apply as an emulsion
Apply the emulsion to the skin after cleansing with pH 4.5-5.5 (pH 7-8) water. Apply a small amount of emulsion to the skin and massage gently. Use only twice a day.

Apply as an oil
Apply the oil to the skin after cleansing with pH 4.5-5.5 (pH 7-8) water. Apply a small amount of oil to the skin and massage gently. Use only twice a day.

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Info-medicament
Indications: For the treatment of the skin.
Contraindications: For the treatment of the skin.
Precautions: For the treatment of the skin.
Warnings: For the treatment of the skin.
Directions: For the treatment of the skin.
Storage: For the treatment of the skin.
Shelf life: For the treatment of the skin.
Batch number: For the treatment of the skin.
Manufacturer: For the treatment of the skin.

PUR 4-in-1 Skin Tint MN4

[illegible]

PUR 4-in-1 Skin Tint MN4

Hydrate, soothe, and help protect your skin with this 40% Water-in-Oil Emulsion. All-in-one moisturizer, sunscreen, and skin care with natural SPF. Sunscreen SPF 15.

Apply as is

Apply liberally to exposed skin using your fingers to blend. Use Layer upon layer of emulsion to achieve the coverage you desire. Reapply as needed. Use also as a pre-sunscreen for protection.

Apply as is

Apply liberally to exposed skin and allow a period of time to blend with skin. Use as a 15-30 min SPF skin care pre-sunscreen. Rehydrate, soothe, and help protect your skin. Reapply as needed. Use also as a pre-sunscreen for protection.

Apply as is

Apply liberally to exposed skin. Use as a pre-sunscreen for protection. Apply liberally to exposed skin. Use as a pre-sunscreen for protection.

Drug Facts (continued)

Indications

Active ingredients

Other ingredients

Warnings

Directions

Other information

Information

Other information

Other information

Other information

Other information

Other information

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Drug Facts (continued)

Indications

Active ingredients

Other ingredients

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zinc oxide lotion

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:67345-0053
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	132 mg in 1 mL

Inactive Ingredients	
Ingredient Name	Strength
TOCOPHEROL (UNII: R0ZB2556P8)	
RETINOL (UNII: G2SH0XKK91)	
ISODODECANE (UNII: A8289P68Y2)	
HELIANTHUS ANNUUS (SUNFLOWER) SEED OIL (UNII: 3W1JG795YI)	
HYDROGENATED SOYBEAN LECITHIN (UNII: H1109Z9J4N)	
CERAMIDE AP (UNII: F1X8L2B00J)	
LACTIC ACID (UNII: 33X04XA5AT)	
BUTYROSPERMUM PARKII (SHEA) BUTTER (UNII: K49155VL9Y)	
POLYGLYCERYL-4 DIISOSTEARATE/POLYHYDROXYSTEARATE/SEBACATE (UNII: 687U3PEB2Y)	
ORYZA SATIVA (RICE) BRAN (UNII: R60QEP13IC)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
POLYMETHYLSILSESQUIOXANE (4.5 MICRONS) (UNII: 59Z907ZB69)	
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)	
ROSMARINUS OFFICINALIS (ROSEMARY) LEAF OIL (UNII: 8LGU7VM393)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
ALLANTOIN (UNII: 344S277G0Z)	
.ALPHA.-BISABOLOL, (+)- (UNII: 105S6I733Z)	
TRILAURETH-4 PHOSPHATE (UNII: M96W2OLL2V)	
BUTYLOCTYL SALICYLATE (UNII: 2EH13UN8D3)	
NIACINAMIDE (UNII: 25X51I8RD4)	
WATER (UNII: 059QF0KO0R)	
DIMETHICONOL/PROPYLSILSESQUIOXANE/SILICATE CROSSPOLYMER (450000000 MW) (UNII: 9KB5R958PB)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
HEXYLENE GLYCOL (UNII: KEH0A3F75J)	
PROPANEDIOL (UNII: 5965N8W85T)	
TETRASODIUM GLUTAMATE DIACETATE (UNII: 5EHL50I4MY)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
ALKYL (C12-15) BENZOATE (UNII: A9EJ3J61HQ)	
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	

LAURYL PEG-8 DIMETHICONE (300 CPS) (UNII: ELL2U7K8T8)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
FERROSO FERRIC OXIDE (UNII: XM0M87F357)	
OCTYLDODECYL NEOPENTANOATE (UNII: X8725R883T)	
CAPRYLYL METHICONE (UNII: Q95M2P1KJL)	

Product Characteristics

Color	brown	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67345-0053-0	1 in 1 CARTON	06/19/2025	
1	NDC:67345-0053-1	30 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	06/19/2025	

Labeler - PUR Minerals (114450443)

Revised: 9/2025

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